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(54) Title: CRYSTALLINE FORMS VI AND VII OF ATORVASTATIN CALCIUM

(57) Abstract: The present invention relates to novel crystalline forms of atorvastatin calcium and to the methods of their production and isolation. More specifically, the present invention relates to novel Forms VI and VII of R-(R', R'')-2-(4-fluorophenyl)-B, d dihydroxy-5(1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-1-H-pyrrole-1-heptanoic acid calcium salt and hydrates thereof and to methods of their preparation.

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## CRYSTALLINE FORMS VI AND VII OF ATORVASTATIN CALCIUM

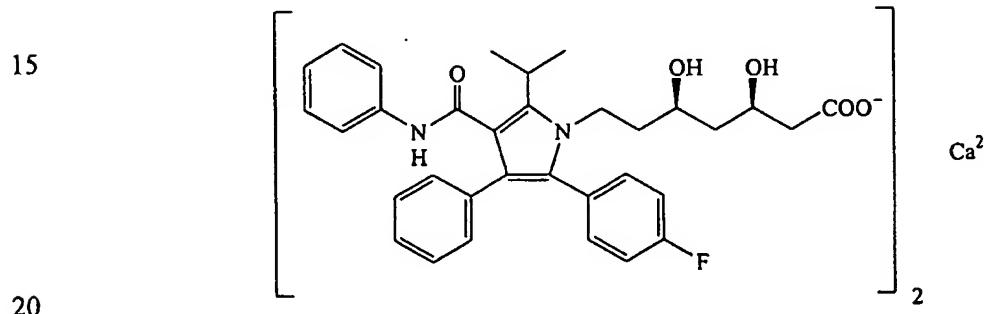
FIELD OF THE INVENTION

The present invention relates to novel crystalline forms of Atorvastatin calcium and to the methods of their production and isolation.

More specifically, the present invention relates to novel Forms VI and VII of R-(R<sup>\*,</sup> R<sup>\*)]-2-(4-fluorophenyl)-β, δ-dihydroxy-5-(1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-1-H-pyrrole-1-heptanoic acid calcium salt and hydrates thereof and to methods of their preparation.</sup>

BACKGROUND OF THE INVENTION

Chemically Atorvastatin is R-(R<sup>\*,</sup> R<sup>\*)]-2-(4-fluorophenyl)-β, δ-dihydroxy-5-(1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-1H-pyrrole-1-heptanoic acid. Atorvastatin is marketed as the hemi calcium salt trihydrate under the name LIPITOR by Warner Lambert Co and may be represented by Formula 1.</sup>



Atorvastatin is a member of the class of drugs called statins. Statin drugs are currently the most therapeutically effective drugs available for reducing low density lipoprotein (LDL) particle concentration in the blood stream of patients at risk for cardiovascular disease. A high level of LDL in to bloodstream 25 has been linked to the formation of coronary lesions, which obstruct the flow of blood and can rupture and promote thrombosis; Goodman Gilman, *The Pharmacological Basis of Therapeutics* 879 (9<sup>th</sup> ed. 1996). Reducing plasma LDL levels has been shown to reduce the risk of clinical events in patients with cardiovascular disease and patients who are free of cardiovascular disease but who have 30 hypercholesterolemia; Scandinavian Simvastatin Survival Study Group, 1994; Lipid Research Clinics Program, 1984a, 1984b.

U.S. Patent 5,929,156 to Warner-Lambert Company, discloses crystalline Form I Atorvastatin hydrate, crystalline Form II Atorvastatin and hydrates thereof and crystalline Form IV Atorvastatin and hydrates thereof, which

are useful as inhibitors of enzyme 3-hydroxy-3methylglutaryl-coenzyme A reductase (HMG-CoA reductase) and are hence useful hypolipidemic and hypocholesterolemic agents.

U.S. Patent 6,121,461 also to Warner-Lambert Company, discloses 5 crystalline Form III Atorvastatin hydrate, which is also a useful hypolipidemic and hypocholesterolemic agent.

WO 01/36384 A1 to Teva Pharmaceutical Industries Ltd discloses Form V Atorvastatin calcium and hydrates thereof; its preparation and a pharmaceutical composition thereof.

10 A process for the preparation of hydrated and anhydrous amorphous Atorvastatin is disclosed in U.S. Patent 6,087,511 also to Warner-Lambert Company.

WO 01/28999 A1 to Egis Gyogyszergyarrt discloses a process for the preparation of amorphous Atorvastatin calcium.

15 It is also noteworthy to point out that U.S. Patent No. 5,969,156 designates the Atorvastatin formed by prior art process (viz United States Patents 5,003,080; 5,097,045; 5,103,024; 5,124,482; 5,149,837; 5,155,251; 5,216,174; .. 5,245,047; 5,248,793; 5,280,126; 5,397,792; and 5,342,952) as amorphous Atorvastatin which has unsuitable filtration and drying characteristics for large 20 scale production and which must be protected from light, heat, oxygen and moisture (column 1; lines 62-65).

25 Atorvastatin is prepared as its calcium salt, which is desirable since it enables Atorvastatin to be conveniently formulated for oral administration. There is hence a need to produce Atorvastatin calcium in a pure and crystalline form to enable formulations to meet exacting pharmaceutical requirements and specifications.

Furthermore, the process by which the crystalline form of Atorvastatin calcium is produced needs to be one, which is amenable to large-scale 30 production. Additionally, it is desirable that the product should be in a form that is readily filterable and easily dried. Finally, it is economically desirable that the product be stable for extended periods of time without the need for specialized storage conditions.

The main aspect of the present invention is to provide novel crystalline forms of Atorvastatin calcium and hydrates thereof, and to a method for

their preparation.

Another aspect of the present invention is that, the novel crystalline forms of Atorvastatin calcium and hydrates thereof are obtained in high purity.

The generally preferred HPLC purity of crystalline Form VI and VII of

5 Atorvastatin calcium and hydrates thereof, of the present invention is greater than 99.0% more preferably greater than 99.5%. Most pharmaceutical formulation processes are facilitated by use of the active materials that are free flowing high melting solids. The novel crystalline forms of Atorvastatin calcium of the present invention are high melting solids, very suited for formulation. The residual  
10 solvents associated with the novel forms, Form VI and Form VII are also very well within permissible limits and that again renders the novel crystalline forms suited for formulations.

#### SUMMARY OF THE INVENTION

Accordingly, the present invention is directed to novel crystalline  
15 forms of Atorvastatin calcium and hydrates thereof. These crystalline forms of Atorvastatin calcium are designated as Form VI and Form VII for convenience.

The present invention further provides a process for the preparation of novel crystalline Form VI and Form VII of Atorvastatin calcium and hydrates thereof, which is a commercially viable process well suited for industrial scale up.

#### BRIEF DESCRIPTION OF ACCOMPANYING DRAWINGS

Fig 1 is a characteristic X ray powder diffractogram of novel crystalline Atorvastatin calcium Form VI. Vertical axis: Intensity (CPS); Horizontal axis: Two Theta (degrees). The significant d values (A°) obtained are 22.52, 19.44, 11.84, 11.23, 9.58, and 4.69.

25 Fig 2 is a characteristic is an X ray powder diffractogram of novel crystalline Atorvastatin calcium Form VII. Vertical axis: Intensity (CPS); Horizontal axis: Two Theta (degrees). The significant d values (A°) obtained are 19.36, 11.80 9.60, 4.75, 4.69 and 4.39.

#### DETAILED DESCRIPTION OF THE INVENTION

30 The present invention is directed to novel crystalline forms of Atorvastatin calcium and hydrates thereof. More particularly, the hydrates contain 1 to 4 moles of water. The crystalline Form VI and Form VII of the present invention may be characterized by their X Ray powder diffraction. Thus the X-Ray diffraction patterns of Form VI and VII of Atorvastatin calcium and their

hydrates were measured on a Rigaku D/Max 2200 Powder Diffractometer with Cu Radiation source. Crystalline Form VI has X-ray powder diffraction pattern essentially as shown in the Table 1. The X-ray powder diffraction pattern is expressed in terms of the  $2\theta$ , d-spacings, and relative intensities > 15 %.

	$2\theta$	d-spacings	Intensity, $I/I_o$ , %
5	3.92	22.52	40
	4.54	19.44	17
	7.46	11.84	100
	7.86	11.23	21
	9.22	9.58	19
	18.9	4.69	55

Table 2 lists the  $2\theta$ , d-spacings, and relative intensities > 15 % for crystalline Form VII of Atorvastatin calcium.

	$2\theta$	d-spacings	Intensity, $I/I_o$ , %
15	4.56	19.36	21
	7.48	11.80	100
	9.20	9.60	21
	18.64	4.75	15
	18.88	4.69	24
	20.20	4.39	17

The present invention is also directed to processes for preparation of novel crystalline forms of Atorvastatin calcium and hydrates thereof.

The present invention hence, provides a process for the preparation of crystalline Form VI of Atorvastatin calcium, which comprises:

25 a. heating a mixture of  $[R-(R^*, R^*)]-2-(4\text{-fluorophenyl})-\beta$ ,  $\delta$ -dihydroxy-5- 1-methyl ethyl-3-phenyl-4-[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tertiary butyl ester (hereinafter referred to as "ester" or "active ingredient" for convenience), acetonitrile and sodium hydroxide flakes to about 25-60°C;

30 b. maintaining the reaction mixture of step a) at 25-60°C for about 3-9 hours preferably 6 hours;

c. adding to the above reaction mixture an aqueous solution of

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calcium acetate hemihydrate;

- d. further stirring the reaction mixture at 30-50°C for about 30 minutes to 2 hours, preferably 45 minutes;
- e. filtering the reaction solution obtained in step d) through 5 hyflow bed;
- f. distilling the solvent from reaction solution of step e) to yield a residue;
- g. suspending the residue of step f) in a mixture of aliphatic nitrile solvent selected from acetonitrile and propionitrile; and water, in a ratio of 10 1:0.1-2, such that the volume of the mixture of solvent- water is 18-40 times the weight of ester in step a); (The ratio of active ingredient to the mixture of solvent and water is 1:18-40 (wt/vol)).
- h. refluxing the reaction mixture obtained in step g) for 10-18 hours preferably 12-14 hours; and
- i. isolating the crystalline Form VI of Atorvastatin calcium, obtained in step h) by filtration or other conventional methods known in the art.

In step a) of the process an ester with an alkyl group of 1-10 carbon atoms, allyl or benzyl group can be used in place of the tertiary butyl ester and another nitrile such as propionitrile can be used in place of acetonitrile. The ratio 20 of solvent to active ingredient in step a) is 16 times the active ingredient (wt:volume) (gm:ml).

In step a) the molar ratio of active ingredient to base is 1:1-1.5 preferably 1:1.15.

In step a) other alkali hydroxides can be used in place of sodium 25 hydroxide. The alkali hydroxides including the sodium hydroxide may be in any form and the form is not limited to flakes.

The following organic and inorganic salts of calcium may be used in place of calcium acetate hemihydrate:

Organic salts such as carboxylates and sulphonates. The 30 carboxylates may be selected from acetate, propionate, butyrate, tartarate; aryl carboxylates like benzoate and phthalate as well as higher carboxylates like Stearate or dodecanoate. Also included are succinate and ascorborate.

Sulphonates may be selected from lower alkyl and aryl sulfonates like calcium methane sulfonates, calcium benzene sulfonates and calcium para

toluenesulfonates.

Inorganic salts of calcium may be selected from calcium chloride, fluoride, bromide, iodide and calcium borate and tetra fluoro borate, calcium carbonate, mono, di and tri basic calcium phosphate, calcium sulfate and calcium hydroxide and hydrates thereof.

The molar ratio of active ingredient to calcium acetate hemihydrate or calcium salt is 1:0.5-0.7 preferably 1:0.6.

The present invention also provides a process for the preparation of crystalline Form VII of Atorvastatin calcium, which comprises:

- 10 a. suspending residue (prepared as per steps a-f for crystalline form VI) in a mixture of water and organic solvent such as an amide solvent such as dimethyl formamide or an aliphatic nitrile solvent selected from acetonitrile or propionitrile; in a ratio of organic solvent to water 1:0.1-5 (vol/vol) such that the volume of the mixture of solvent- water is 5 to less than 10 times, the weight of initial ester used in the preparation of crystalline form VI (vol:wt) (ml/gm);
- 15 b. refluxing the reaction mixture obtained in step a) for 10 minutes to 1 hour preferably 30 minutes;
- 20 c. subsequently, adding a second mixture of organic solvent: water (1:0.01-1) (vol/vol); such that the volume of the mixture of solvent- water is 5-10 times the weight of the initial ester; and refluxing the reaction mixture for 10 minutes to 1 hours, preferably 30 minutes;
- 25 d. finally, adding a third mixture of organic solvent: water (1:0.01-1) (vol/vol); such that the volume of mixture of solvent- water is 5-10 times the weight of the initial ester; and refluxing the reaction mixture for 1 hour to 3 hours, preferably 1 hour;
- 30 e. cooling the reaction mixture of step d) to 0-10°C preferably below 5°C; and
- f. isolating the crystalline Form VII of Atorvastatin calcium, obtained in step e) by filtration or other conventional methods known to art.

The organic solvents used in steps c) and d) for the preparation of crystalline Form VII of Atorvastatin calcium include amide solvents such as dimethyl formamide or aliphatic nitrile solvent selected from acetonitrile and propionitrile. The same solvent used in step a) is used in steps c) and d).

The present invention hence provides novel crystalline forms of

Atorvastatin calcium and hydrates thereof, and to a method for their preparation, which is amenable to large-scale production.

The novel crystalline forms of Atorvastatin calcium of the present invention are readily filterable and easily dried.

5 Moreover, the HPLC purity of novel crystalline Form VI and VII of Atorvastatin calcium and hydrates thereof, of the present invention is greater than 99.0% preferably greater than 99.5%.

10 The crystalline forms of Atorvastatin calcium of the present invention are also high melting solids with residual solvents within permissible limits and are very well suited for formulation. The crystalline Form VI of Atorvastatin calcium hydrate may contain 1 to 4 moles of water, preferably 3 moles of water. The crystalline Form VI of Atorvastatin calcium hydrate may contain 1 to 5 moles of water, preferably 3 moles of water.

#### EXAMPLES

15 The present invention is illustrated by the following examples, which are not intended to limit the effective scope of the invention.

##### Example 1

###### Preparation of crystalline Form VI of Atorvastatin Calcium

20 A mixture of [R-(R\*, R\*)]-2-(4-fluoro phenyl) $\beta$ ,  $\delta$ -dihydroxy-5-[(1-methyl ethyl)- 3-phenyl-4[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tert. butyl ester (25.0g), acetonitrile (400ml) water (62 ml) and sodium hydroxide flakes (1.88g) are heated to about 25-55°C and maintained at the same temperature for about 4 and a half hours. To the reaction mixture is then added an aqueous solution of calcium acetate hemihydrate (4.0g in 41.6 ml of water) and 25 stirred at 30-50°C for about 1 hour. Subsequently, the solution obtained is filtered through hyflow bed and washed with acetonitrile (125 ml). The solvent is then distilled completely to yield residue.

30 To the residue thus obtained, a mixture of acetonitrile: water (1:1) (500 ml) is added and the reaction mixture maintained at reflux for about 13 hours. The separated solid is then filtered at 70°C and washed with a mixture of acetonitrile: water (1:1) (50 ml). The resultant solid is dried at 60-70°C to render desired crystalline Form-VI of Atorvastatin Calcium.

HPLC Purity: 99.71%

Example 2Preparation of crystalline Form VII of Atorvastatin Calcium

A mixture of [R-(R\*, R\*)]-2-(4-fluoro phenyl)- $\beta$ ,  $\delta$ -dihydroxy-5-[(1-methyl ethyl)-3-phenyl-4[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tert. butyl ester (25.0g), acetonitrile (400 ml) and sodium hydroxide flakes (1.88g) are heated to about 30-45°C and maintained at the same temperature for about 6 hours. To the reaction mixture is then added an aqueous solution of calcium acetate hemihydrate (4.0g in 41.6 ml of water) and stirred at 30-50°C for about 50 minutes. Subsequently, the solution obtained is filtered through hyflow bed and washed with acetonitrile (125 ml). The solvent is then distilled completely to yield residue. To the residue thus obtained, a mixture of acetonitrile: water (1:1; 150 ml) was added and the reaction mixture maintained at reflux for about 15-20 minutes. Upon completion of this step, a second mixture of acetonitrile: water (1:1; 150ml) is added to resultant slurry, and again the reaction mixture maintained at reflux about 30 minutes. Finally a third mixture of acetonitrile water (1:1; 150ml) is added to resultant slurry, and the reaction mixture maintained at reflux for about 1 hour. The reaction mixture then cooled to 0-10°C, filtered and dried at 50-60°C to get the desired crystalline Form VII of Atorvastatin Calcium.

HPLC Purity: 99.81%

Example 3

A mixture of [R-(R\*, R\*)]-2-(4-fluoro phenyl)- $\beta$ ,  $\delta$ -dihydroxy-5-[(1-methyl ethyl)-3-phenyl-4[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tert. butyl ester (10.0g), acetonitrile(80 ml) and sodium hydroxide flakes (0.75g) in water (25.0 ml) are heated to about 35-45°C and maintained at the same temperature for about 1-2 hours.

To the reaction mixture is then slowly added an aqueous solution of calcium acetate hemihydrate (1.6 g in 16.0 ml of water) at 35-45°C for about 30-60 minutes. After another 15-20 minutes, the temperature is raised to 50-60°C and the solution obtained is filtered through hyflow bed and washed with acetonitrile (10.0 ml). The solvent is then distilled completely to yield residue. To the residue thus obtained, is added acetonitrile (60 ml) and the temperature is raised to 50-60°C and the solution obtained is again filtered through hyflow bed and washed with acetonitrile (10.0 ml). To the filtrate, water (120.0 ml) was added and the reaction mixture maintained at reflux about 60-90 minutes.

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The reaction mixture is then cooled to 0-10°C, filtered and dried at 50-60°C for 10-12 hours to get the desired crystalline Form VII of Atorvastatin Calcium.

Yield 7 to 9 gm

5 HPLC Purity 99.7%

CLAIMS

1. A crystalline Form VI of R-(R<sup>1</sup>,R<sup>2</sup>)-2-(4-fluorophenyl)- $\beta$ ,  
 $\delta$ -dihydroxy-5- (1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-H-pyrrole-1-heptanoic acid hemi calcium salt (Atorvastatin calcium) or hydrates thereof.
- 5 2. A crystalline Form VI of Atorvastatin calcium hydrate as claimed in claim 1, which is characterized by the following X-ray powder diffraction pattern (d values in A°): 22.52, 19.44, 11.84, 11.23, 9.58, and 4.69.
3. A process for preparing Form VI of Atorvastatin calcium or hydrates thereof, which comprises:
  - 10 a. heating a mixture of [R-(R<sup>1</sup>, R<sup>2</sup>)-2-(4-fluoro phenyl)- $\beta$ ,  $\delta$ -dihydroxy-5- [(1-methyl ethyl)-3-phenyl- 4-[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tert. butyl ester, acetonitrile and sodium hydroxide flakes to about 25-60°C;
  - b. maintaining the reaction mixture of step a) at 25-60°C for about 3
  - 15 to 9 hours preferably 6 hours;
  - c. adding to the above reaction mixture an aqueous solution of calcium acetate hemihydrate;
  - d. further stirring the reaction mixture at 30-50°C for about 1-2 hours preferably 45 minutes;
  - 20 e. filtering the reaction solution obtained in step d) through hyflow bed;
  - f. distilling the solvent from reaction solution of step e) to a yield residue;
  - 25 g. suspending the residue of step f) in a mixture of water and aliphatic nitrile solvent selected from acetonitrile and propionitrile;
  - h. refluxing the reaction mixture obtained in step g) for 10-18 hours, preferably 12-14 hours; and
  - i. isolating the crystalline Form VI of Atorvastatin calcium, obtained in step h).
- 30 4. The process as claimed in claim 3, wherein in step g) the ratio of nitrile solvent to water is 1:0.1-2.
5. The process as claimed in claim 3, wherein in step g) the volume of the mixture of solvent and water is 18-40 times the weight of the ester added in step a).

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6. The process as claimed in claim 3, wherein in step a) the molar ratio of  $[R-(R', R'')]-2-(4\text{-fluoro phenyl})-\beta, \delta\text{-dihydroxy-5-[(1-methyl ethyl)-3-phenyl-4-[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, tert butyl ester to sodium hydroxide is 1:1-1.5, preferably 1:1.15.}$
- 5 7. The crystalline Form VI of Atorvastatin calcium hydrate according to any one of claims 1 to 2, which contains from 1 to 4 moles of water.
8. The crystalline Form VI of Atorvastatin calcium hydrate according to any one of claims 1 to 2, which contains from 3 moles of water.
9. A crystalline Form VII of  $R-(R', R'')]-2-(4\text{-fluorophenyl})\beta, \delta\text{-dihydroxy-5- (1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-1H-pyrrole-1-heptanoic acid hemi calcium salt (Atorvastatin calcium) or hydrates thereof.}$
- 10 10. The crystalline Form VII of Atorvastatin calcium hydrate, as claimed in claim 9, which is characterized by the following X-ray powder diffraction pattern (d values in  $\text{\AA}$ ): 19.36, 11.80, 9.60, 4.75, 4.69 and 4.39.
- 15 11. A process for preparing Form VII of Atorvastatin calcium and hydrates thereof, which comprises:
  - a) suspending a residue prepared according to steps a) to f) of claim 3 in a mixture of water and organic solvent selected from an amide solvent or an aliphatic nitrile solvent;
  - 20 b) refluxing the mixture obtained in step a) for 10 minutes to 1 hours preferably 30 minutes;
  - c) adding a second mixture of the water and the organic solvent of step a) to the mixture of step b) and refluxing the reaction mixture for 10 minutes to 1 hours preferably 30 minutes;
  - 25 d) adding a third mixture of the water and the organic solvent of step a) to the mixture of step c) and refluxing the reaction mixture for 1 - 3 hours preferably 1 hour;
  - e) cooling the reaction mixture of step d) to 0-10°C preferably below 5°C; and
  - 30 f) isolating the crystalline Form VII of Atorvastatin calcium obtained in step e).
12. The process as claimed in claim 11, wherein the ratio of organic solvent to water in step a) is 1:0.1-5 (vol/vol).
13. The process as claimed in claim 11, wherein in step a) the volume of

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the mixture of organic solvent and water is 5 to less than 10 times the weight of the initial ester added in step a).

14. The process as claimed in claim 11, wherein in step c) the volume of the mixture of organic solvent and water is 5-10 times the initial ester.

5 15. The process as claimed in any one of claims 11 to 13, wherein the organic solvent is dimethyl formamide.

16. The process as claimed in any one of claims 11 to 13, wherein the aliphatic nitrile is selected from acetonitrile or propionitrile.

17. The crystalline Form VI of Atorvastatin calcium hydrate according to 10 any one of claims 9 to 10, which contains 1 to 5 moles of water.

18. The crystalline Form VI of Atorvastatin calcium hydrate according to any one of claims 9 to 10, which contains 3 moles of water

19. A process for preparing Form VI of Atorvastatin calcium or hydrates thereof, which comprises:

15 a. heating a mixture of [R-(R<sup>1</sup>, R<sup>2</sup>)]-2-(4-fluoro phenyl)-β, δ-dihydroxy-5- [(1-methyl ethyl)-3-phenyl- 4-[(phenyl amino)-carbonyl]-1H-pyrrole-1-heptanoic acid, a compound selected from C<sub>1</sub>-C<sub>10</sub> alkyl ester, allyl ester or benzyl ester; nitrile and alkali hydroxide to about 25-60°C;

20 b. maintaining the reaction mixture of step a) at 25-60°C for about 3 to 9 hours preferably 6 hours;

c. adding to the reaction mixture of step b) an aqueous solution of a calcium salt;

d. further stirring the reaction mixture at 30-50°C for about 1-2 hours preferably 45 minutes;

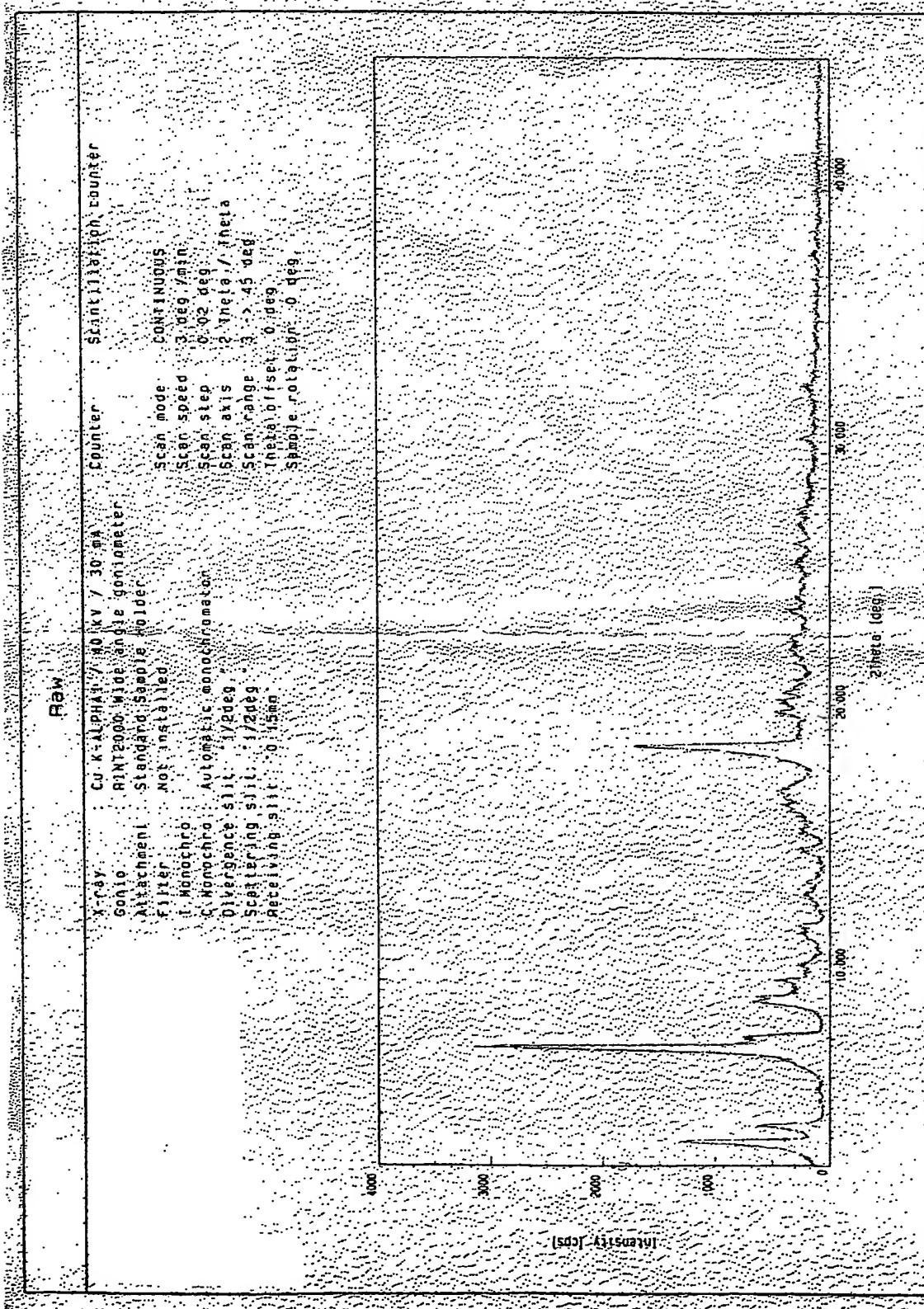
25 e. filtering the reaction solution obtained in step d) through hyflow bed;

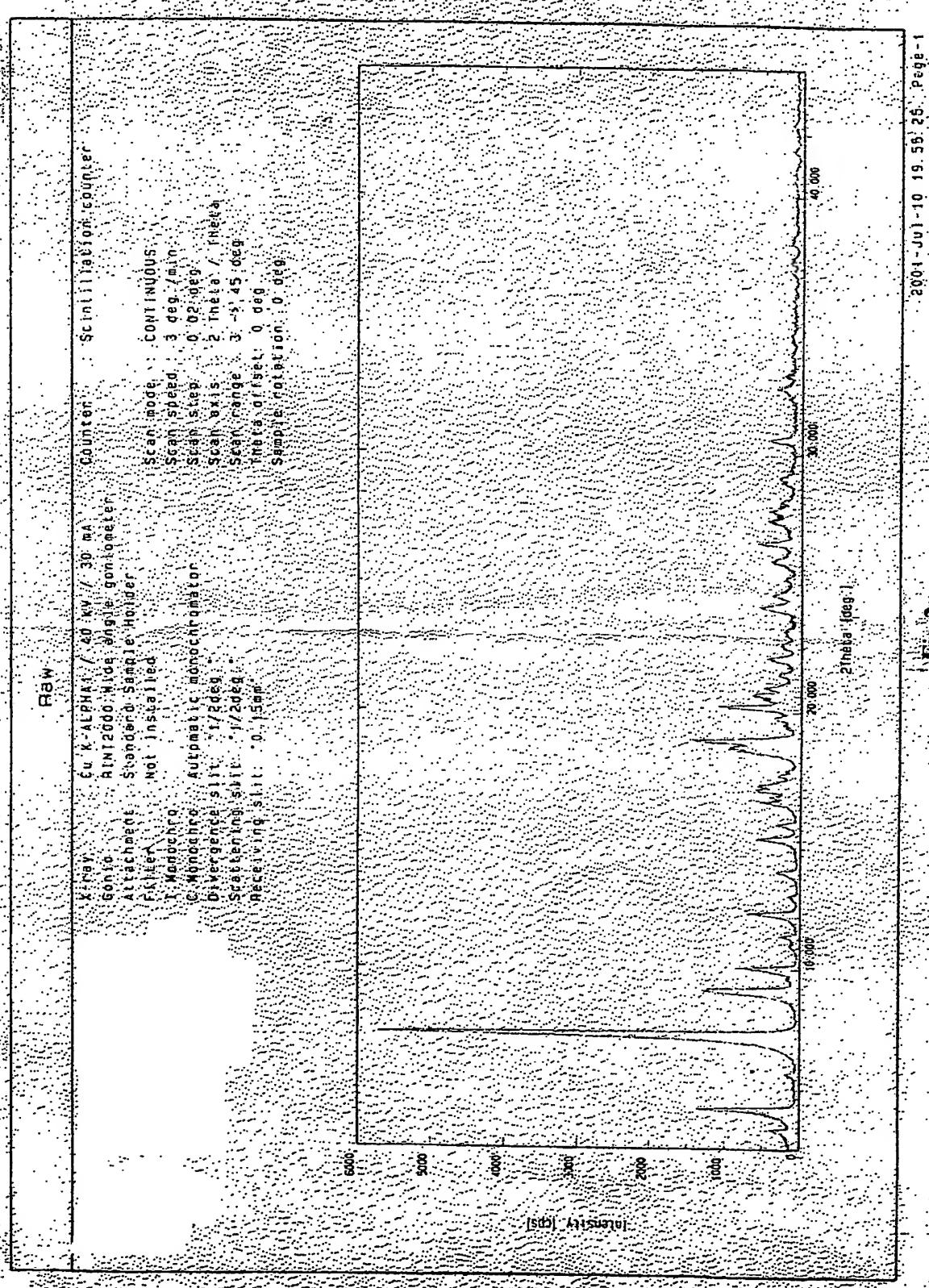
f. distilling the solvent from reaction solution of step e) to yield a residue;

g. suspending the residue of step f) in a mixture of aliphatic nitrile 30 solvent selected from acetonitrile or propionitrile and water;

h. refluxing the mixture of step g) for 10-18 hours; preferably 12-14 hours; and

i. isolating the crystalline Form VI of Atorvastatin calcium.





## INTERNATIONAL SEARCH REPORT

In	Application No
PCT/US 02/00431	

A. CLASSIFICATION OF SUBJECT MATTER
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According to International Patent Classification (IPC) or to both national classification and IPC
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B. FIELDS SEARCHED
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Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
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Electronic data base consulted during the international search (name of data base and, where practical, search terms used)
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EPO-Internal, WPI Data, PAJ, CHEM ABS Data
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C. DOCUMENTS CONSIDERED TO BE RELEVANT
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Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 97 03958 A (WARNER LAMBERT CO ;MCKENZIE ANN T (US)) 6 February 1997 (1997-02-06) cited in the application claims 1,2 ---	1-19
X	WO 97 03959 A (WARNER LAMBERT CO ;BRIGGS CHRISTOPHER A (US); JENNINGS REX ALLEN () 6 February 1997 (1997-02-06) cited in the application claims 1-7 ---	1-19
X	WO 01 36384 A (TEVA PHARMA ;AYALON ARI (IL); NIDDAM VALERIE (IL); ROYTBLAT SOFIA) 25 May 2001 (2001-05-25) claims 1-3 ---	1-19 -/-

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- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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Date of the actual completion of the international search
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16 October 2002
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Date of mailing of the international search report
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## INTERNATIONAL SEARCH REPORT

Int	al Application No
PCT/US 02/00431	

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E	WO 02 43732 A (ISHAI ETI ;SAMBURSKY GUY (IL); TEVA PHARMA (IL); ARONHIME JUDITH ()) 6 June 2002 (2002-06-06) claims 1-146 ---	1-19
E	WO 02 057229 A (GANESH SAMBASIVAM ;MATHEW JOY (IN); BIOCON INDIA LTD (IN)) 25 July 2002 (2002-07-25) claim 1 ---	1-19
E	WO 02 41834 A (TEVA PHARMA ;TEVA PHARMACEUTICALS USA INC (US)) 30 May 2002 (2002-05-30) claims 1-4 ---	1-19
E	WO 02 051804 A (SCHOENING KAI-UWE ;SZELAGIEWICZ MARTIN (CH); VAN DER SCHAAF PAUL A) 4 July 2002 (2002-07-04) claims 1-8 ---	1-19
E	WO 02 43667 A (ISHAI ETI ;TEVA PHARMA (IL); NIDDAM VALERIE (IL); LIDOR HADAS RAMY) 6 June 2002 (2002-06-06) scheme 5 claim 14 -----	1-19

## INTERNATIONAL SEARCH REPORT

Information on patent family members

Int'l Application No

PCT/US 02/00431

Patent document cited in search report	Publication date	Patent family member(s)	Publication date	
WO 9703958	A 06-02-1997	AT 207465 T AU 725368 B2 AU 6484196 A BG 63629 B1 BG 102186 A BR 9610567 A CA 2220458 A1 CN 1190957 A , B CZ 9800123 A3 DE 69616358 D1 DK 848704 T3 EE 9800016 A EP 0848704 A1 ES 2166456 T3 HR 960313 A1 HU 9901687 A2 IL 122162 A JP 11509229 T JP 3296563 B2 NO 980208 A NZ 312906 A PL 324532 A1 SK 5998 A3 TW 401399 B WO 9703958 A1 US 6121461 A		15-11-2001 12-10-2000 18-02-1997 31-07-2002 30-10-1998 06-07-1999 06-02-1997 19-08-1998 17-06-1998 29-11-2001 04-02-2002 17-08-1998 24-06-1998 16-04-2002 30-04-1998 28-10-1999 14-07-1999 17-08-1999 02-07-2002 16-01-1998 22-12-2000 08-06-1998 06-05-1998 11-08-2000 06-02-1997 19-09-2000
WO 9703959	A 06-02-1997	AT 208375 T AU 725424 B2 AU 6484296 A BG 63630 B1 BG 102187 A BR 9609872 A CA 2220018 A1 CN 1190955 A , B CZ 9800121 A3 DE 69616808 D1 DE 69616808 T2 DK 848705 T3 EE 9800015 A EP 1148049 A1 EP 0848705 A1 ES 2167587 T3 HR 960339 A1 HU 9900678 A2 IL 122118 A JP 11509230 T JP 3296564 B2 NO 980207 A NZ 312907 A PL 324496 A1 PT 848705 T SI 848705 T1 SK 6298 A3 WO 9703959 A1 US 5969156 A	15-11-2001 12-10-2000 18-02-1997 31-07-2002 30-10-1998 23-03-1999 06-02-1997 19-08-1998 14-10-1998 13-12-2001 29-05-2002 04-02-2002 17-08-1998 24-10-2001 24-06-1998 16-05-2002 30-04-1998 28-07-1999 14-07-1999 17-08-1999 02-07-2002 16-01-1998 22-12-2000 25-05-1998 28-02-2002 30-04-2002 07-10-1998 06-02-1997 19-10-1999	
WO 0136384	A 25-05-2001	AU 1617301 A	30-05-2001	

## INTERNATIONAL SEARCH REPORT

Information on patent family members

In **International Application No****PCT/US 02/00431**

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 0136384	A	EP WO	1235799 A1 0136384 A1	04-09-2002 25-05-2001
WO 0243732	A 06-06-2002	AU WO AU WO US	1792702 A 0243732 A1 3289102 A 0243667 A2 2002099224 A1	11-06-2002 06-06-2002 11-06-2002 06-06-2002 25-07-2002
WO 02057229	A 25-07-2002	WO WO	02057229 A1 02057274 A1	25-07-2002 25-07-2002
WO 0241834	A 30-05-2002	AU WO US	4150602 A 0241834 A2 2002115709 A1	03-06-2002 30-05-2002 22-08-2002
WO 02051804	A 04-07-2002	WO	02051804 A1	04-07-2002
WO 0243667	A 06-06-2002	AU WO US AU WO	3289102 A 0243667 A2 2002099224 A1 1792702 A 0243732 A1	11-06-2002 06-06-2002 25-07-2002 11-06-2002 06-06-2002